510(k) Summary
(21 CFR Part 807.92)

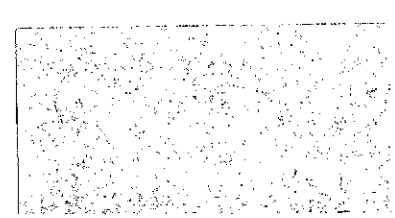
A. Submitter Information

Submitter's Name:	Theken Spine, LLC
Address:	1800 Triplett Blvd Akron, Ohio 44306
Telephone Number:	330-475-8600
Fax Number:	330-773-7697
Contact Person:	Dale Davison
Date Prepared:	4/29/2009

JUN 10 2009

B. Device Information

Trade Name:	Coral™ Spinal System
Common Name:	Pedicle Screw Spinal System
Classification:	**MNI** 888.3070 – Pedicle Screw Spinal System **MNH** 888.3070 – Pedicle Screw Spinal System **KWQ** 888.3060 – Spinal Intervertebral Body Fixation Orthosis **KWP** 888.3050 – Spinal Interlaminal Fixation Orthosis **NKB** 888.3070 – Spondylolisthesis Spinal Fixation System
Predicate Device:	Theken Surgical Coral™ Spinal System, K041592 Theken Surgical Coral™ Spinal System, K070962 Theken Surgical Coral™ Spinal System, K081414 K2M Range Spinal System, K080792 Pioneer Surgical Quantum Spinal System, K070551
Device Description:	The purpose of this submission is the addition of cobalt chrome rods to the Coral™ Spinal System. The Coral™ Spinal System components can be rigidly locked together in a variety of configurations to promote fusion for a wide variety of patient anatomies.
Intended Use:	The Coral™ Spinal System is a non-cervical spinal fixation device intended for use as a posterior pedicle screw fixation system, a posterior non-pedicle screw fixation system, or as an anterolateral fixation system. Pedicle screw fixation is limited to skeletally mature patients. The device is indicated as an adjunct to fusion for the following indications: degenerative disc disease (defined as discogenic back pain with degeneration of the disc confirmed by history and radiographic studies), spondylolisthesis, trauma (i.e., fracture or dislocation), spinal stenosis, deformities or curvatures (i.e., scoliosis, kyphosis, and/or lordosis), tumor, pseudoarthrosis, and failed previous fusion.
Material Composition:	Implant grade titanium alloy Ti 6Al-4V (ELI) per ASTM F-136, CP titanium per ASTM F-67 and CoCr Alloy per ASTM F-1537.



C. Substantial Equivalence

Theken Spine believes sufficient evidence exists to reasonably conclude that the additional components are substantially equivalent to the predicate device Coral™ Spinal System (K041592 SE 9/04, K070962 SE 8/07, and K081414 SE 7/08), manufactured by Theken Spine, LLC. This is based on the design concept, the use of established, known materials, feature comparisons, mechanical testing, indications for use, pre-production quality assurance planning and engineering analysis. All implants are used to treat the same conditions, possess the same precautions and contraindications for use, and equivalent potential for complications for the risk of use.

The subject device similarities include:

- The same indications for use
- The same operating principle
- The same biocompatible materials
- Implanted using the same surgical techniques and equipment type
- The same manufacturing environment
- The same sterilization process
- The same packaging configurations

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Theken Spine LLC
% Mr. Dale Davison
Vice President-Engineering
1800 Triplett Boulevard
Akron, Ohio 44306

JUN 10 2009

Re: K091266
Trade/Device Name: Coral™ Spinal System
Regulation Number: 21 CFR 888.3070
Regulation Name: Pedicle Screw Spinal System
Regulatory Class: III
Product Code: NKB, MNI, MNH, KWP, & KWQ
Dated: April 29, 2009
Received: May 13, 2009

Dear Mr. Davison:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing

practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please go to http://www.fda.gov/AboutFDA/CentersOffices/CDRH/CDRHOffices/ucm115809.htm for the Center for Devices and Radiological Health's (CDRH's) Office of Compliance. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to http://www.fda.gov/cdrh/mdr/ for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

Mark N. Melkerson
Division of Surgical, Orthopedic
and Restorative Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known): K091266

The Coral™ Spinal System is a non-cervical spinal fixation device intended for use as a posterior pedicle screw fixation system, a posterior non-pedicle screw fixation system, or as an anterolateral fixation system. Pedicle screw fixation is limited to skeletally mature patients. The device is indicated as an adjunct to fusion for the following indications: degenerative disc disease (defined as discogenic back pain with degeneration of the disc confirmed by history and radiographic studies), spondylolisthesis, trauma (i.e., fracture or dislocation), spinal stenosis, deformities or curvatures (i.e., scoliosis, kyphosis, and/or lordosis), tumor, pseudoarthrosis, and failed previous fusion.

Prescription Use ___X____ AND/OR Over-The-Counter Use _____

(Part 21 CFR 801 Subpart D) (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

for (Division Sign-Off)
Division of Surgical, Orthopedic,
and Restorative Devices

510(k) Number K091266